In the Claims

- 1. (Currently Amended) A process for making neutral or anionic complexes containing sequestered DNA for gene transfer, comprising: forming a stable colloid comprising an aqueous phase having suspended therein a first DNA complex with a cationic surface potential comprising a DNA sequence complexed with a cationic lipid or polymer comprising one or more cationic head groups; and modifying the surface potential of said first DNA complex to form a stable colloid comprising a second DNA complex with a neutral or net anionic surface potential by reacting said cationic head groups with a reagent that reacts with the cationic head group to neutralize the positive charge thereon.
- 2. (Withdrawn) The process of claim 1, wherein the surface potential of said first DNA complex is modified by adding a poly(alkylene oxide) to the aqueous phase of said colloid.
- 3. (Withdrawn) The process of claim 2, wherein said poly(alkylene oxide) is polyetheylene glycol.
- 4. (Withdrawn) The process of claim 1, wherein the surface potential of said first DNA complex is modified by the covalent attachment of poly(alkylene oxides) to the cationic lipid or polymer.
- 5. (Withdrawn) The process of claim 4, wherein said poly(alkylene oxide) is polyethylene glycol.

- 6. (Cancelled)
- 7. (Currently Amended) The process of claim <u>1</u> 6, wherein said cationic lipid or polymer is selected from the group consisting of linear polyamines, branched polyamines and polyamines comprising guanidinium groups.
- 8. (Currently Amended) The process of claim 1 6, wherein said reagent is citraconic anhydride or N-hydroxysuccimide acetate.
- 9. (Currently Amended) The process of claim 1 6, wherein said reagent is an N-hydroxysuccinimide ester of a targeting ligand, so that a targeting ligand is covalently attached to said cationic lipid or polymer that also modifies the surface potential of said first DNA complex.
- 10. (Original) The process of claim 9, wherein said targeting ligand is an amino sugar or peptide.
- 11. (Original) The process of claim 1, wherein said first DNA complex further comprises a targeting ligand covalently attached to said cationic lipid or polymer.
- 12. (Withdrawn) The process of claim 4, wherein said poly(alkylene oxide) is only covalently attached to cationic lipids or polymers on the surface of said first DNA complex.
- 13. (Withdrawn) The process of claim 4, wherein said poly(alkylene oxide) is covalently attached to cationic lipids or polymers on the surface of and in the

interior of said first DNA complex.

- 14. (Currently Amended) The process of claim 1 6, wherein said reagent is only reacted with cationic head groups of cationic lipids or polymers on the surface of said first DNA complex.
- 15. (Currently Amended) The process of claim 1 6, wherein said reagent is reacted with cationic head groups of cationic lipids or polymers on the surface of and in the interior of said first DNA complex.
- 16. (Withdrawn) A stable colloid comprising an aqueous phase having suspended therein a first DNA complex with a cationic surface potential comprising an exogenous therapeutic DNA sequence for delivery in vivo to a patient in need thereof, complexed with a cationic lipid or polymer, wherein said aqueous phase comprises an aqueous solution of a poly(alkylene oxide).
- 17. (Withdrawn) A stable colloid comprising an aqueous phase having suspended therein a first DNA complex with a cationic surface potential comprising an exogenous therapeutic DNA sequence for delivery in vivo to a patient in need thereof, complexed with a cationic lipid or polymer, wherein said surface potential of said first DNA complex is modified by the covalent attachment of poly(alkylene oxides) to the cationic lipid or polymer.
- 18. (Currently Amended) A stable colloid comprising an aqueous phase having suspended therein a first DNA complex with a cationic surface potential comprising an exogenous therapeutic DNA sequence of delivery in vivo to a patient in need thereof, complexed with a cationic lipid or polymer

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comprising one or more cationic head groups modified by reaction with a reagent that <u>reacts</u> with the cationic head group and neutralizes the positive charge thereon.

19. (Currently Amended) A method for gene therapy by delivering in vivo an exogenous therapeutic DNA sequence to a patient in need thereof comprising administering to said patient an effective amount of the colloid of claim 16, 17 or 18.